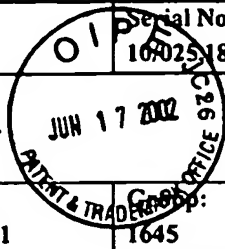


Form PTO-1449 (modified)		Atty. Docket No. IOWA:034US	Serial No. 167025187
List of Patents and Publications for Applicant's  INFORMATION DISCLOSURE STATEMENT  (Use several sheets if necessary)		Applicant Val Sheffield <i>et al.</i>	
		Filing Date: December 18, 2001	
U.S. Patent Documents See Page 1	Foreign Patent Documents See Page 1	Other Art See Page 1	

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## U.S. Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Name	Class	Sub Class	Filing Date of App.

## Foreign Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Country	Class	Sub Class	Translation Yes/No

## Other Art (Including Author, Title, Date Pertinent Pages, Etc.)

Exam. Init.	Ref. Des.	Citation
RT	C1	Bruford <i>et al.</i> , "Linkage mapping in 29 Bardet-Biedl syndrome families confirms loci in chromosomal regions 11q13, 15q22.3-q23, and 16q21," <i>Genomics</i> , 41:93-99, 1997.
	C2	Carmi <i>et al.</i> , "Phenotypic differences among patients with Bardet-Biedl syndrome linked to three different chromosome loci," <i>Am. J. Med. Genet.</i> , 59:199-203, 1995.
	C3	Carmi <i>et al.</i> , "Use of a DNA pooling strategy to identify a human obesity syndrome locus on chromosome 15," <i>Hum. Mol. Genet.</i> , 4:9-13, 1995.
	C4	Farag and Teebi, "Bardet-Biedl and Laurence-Moon syndromes in a mixed Arab population," <i>Clinical Genet.</i> , 33:78-82, 1988.
	C5	Farag and Teebi, "High incidence of Bardet Biedl syndrome among the Bedouin [letter]," <i>Clinical Genet.</i> , 36:463-464, 1989.
/	C6	Katsanis <i>et al.</i> , "Mutations in MKKS cause obesity, retinal dystrophy and renal malformations associated with bardet-biedl syndrome," <i>Nat. Genet.</i> , 26:67-70, 2000.
	C7	Kwitek-Black <i>et al.</i> , "Linkage of Bardet-Biedl syndrome to chromosome 16q and evidence for non-allelic genetic heterogeneity," <i>Nature Genet.</i> , 5:392-396, 1993.
	C8	Leppert <i>et al.</i> , "Bardet-Biedl syndrome is linked to DNA markers on chromosome 11q and is genetically heterogeneous," <i>Nature Genet.</i> , 7:108-112, 1994.
I	C9	Nishimura <i>et al.</i> , "The forkhead transcription factor gene FKHL7 is responsible for glaucoma phenotypes which map to 6p25," <i>Nature Genet.</i> , 19:140-147, 1998.

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### Other Art (Including Author, Title, Date Pertinent Pages, Etc.)

Exam. Init.	Ref. Des.	Citation
RT	C10	Sheffield <i>et al.</i> , "Identification of a Bardet-Biedl syndrome locus on chromosome 3 and evaluation of an efficient approach to homozygosity mapping," <i>Hum. Mol. Genet.</i> , 3(8):1331-1335, 1994.
	C11	Slavotinek <i>et al.</i> , "Mutations in MKKS cause bardet-biedl syndrome," <i>Nature Genet.</i> , 26:15-16, 2000.
	C12	Stone <i>et al.</i> , "Mutation of a gene encoding a putative chaperonin causes McKusick-Kaufman syndrome," <i>Nat. Genet.</i> , 25:79-82, 2000.
	C13	Swiderski <i>et al.</i> , "Expression pattern and in situ localization of the mouse homologue of the human MYCO (GLC1A) gene in adult brain," <i>Mol. Brain Res.</i> , 68:64-72, 1999.
	C14	Woods <i>et al.</i> , "Genetic heterogeneity of Bardet-Biedl syndrome in a distinct Canadian population: evidence for a fifth locus," <i>Genomics</i> , 55:2-9, 1999.
	C15	Young <i>et al.</i> , "A fifth locus for Bardet-Biedl syndrome maps to chromosome 2q31," <i>Am. J. Hum. Genet.</i> , 64:900-904, 1999.

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